

Please amend claims 1, 2, 6, 7, 8, 9, 12, 13, 14 and 15 as follows:

1. (Twice Amended) A peptide of the formula: $R^1-X^1-X^2-R^2$

wherein X^1 is phenyl alanine

Sub E1 } X^2 is any amino acid residue;

Sub E1 } R^1 is NH_2 - or an amino acid sequence $X^3-X^4-X^5$

D2 } wherein X^3 is an aliphatic amino acid residue having a side chain hydroxyl group
and X^4 and X^5 are the same or different and are any amino acid residue and wherein R^2 is a
sequence of 1 to 3 amino acid residues which are the same or different and are aliphatic
amino acid residues provided that the peptide is not Phe-Glu-Gly, Phe-Ala-Gly or Phe-Ala-
Gly-Gly.

2. (Twice Amended) [The] A peptide of [claim 1] the formula: $R^1-X^1-X^2-R^2$

wherein X^2 is Glu or Ala;

Sub E2 } R^2 is Gly-Gly;

Sub E2 } R^1 is $X^3-X^4-X^5$ wherein

Sub E2 } X^4 is Asp or Ala and

Sub E2 } X^5 is Ile or Ala

6. (Amended) The peptide of claim [3] 2 having an amino acid sequence

D2 } selected from the group consisting of:

(a) Phe-Glu-Gly-Gly-Gly (Sequence ID NO:9);

Sub E2 cont

[(b) Phe-Glu-Gly; and]
(b) Phe-Glu-Gly-Gly;
(c) Phe-Ala-Gly-Gly-Gly; and
[(c)](d) Phe-Glu-Sarcosine.

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7. (Amended) [The A peptide of [claim 1] the formula: R¹-X¹-X²-R² wherein X¹ is phenyl alanine;
X² is any amino acid residue;
R¹ is NH₂- or an amino acid sequence X³-X⁴-X⁵
wherein X³ is an aliphatic amino acid residue having a side chain hydroxyl group
and X⁴ and X⁵ are the same or different and are any amino acid residue and wherein R² is a
sequence of 1 to 3 amino acid residues which are the same or different and are selected
from the group consisting of [glycine,] sarcosine, azetidine, nipecotic acid and pipecotic
acid.

2 *1*
8. (Amended) The peptide of claim [3] wherein [R² is a sequence of 1 to 3
amino acid residues which are the same or different and are selected from the group
consisting of glycine, sarcosine, azetidine, nipecotic acid and pipecotic acid] R¹ is NH₂-
and X² is Glu or Ala.

CHF21
9. (Amended) The peptide of claim [1]2 wherein at least one amino acid is a D
amino acid.

Sub E3
10. (Amended) The peptide of claim [4]6 wherein Phe and Glue or Ala are D
amino acids.

12. (Amended) A pharmaceutical composition comprising a peptide of [claim 1]
the formula: $R^1-X^1-X^2-R^2$

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wherein X^1 is phenyl alanine

X^2 is any amino acid residue,

R^1 is NH_2- or an amino acid sequence $X^3-X^4-X^5$

wherein X^3 is an aliphatic amino acid residue having a side chain hydroxyl group
and X^4 and X^5 are the same or different and are any amino acid residue and wherein R^2 is a
sequence of 1 to 3 amino acid residues which are the same or different and are aliphatic
amino acid residues and a pharmaceutically acceptable carrier.

CHF31
13. (Amended) A method for treating or preventing SIRS-induced hypotension
in a mammal comprising administering to the mammal an effective amount of a peptide of
[claim 1] the formula: $R^1-X^1-X^2-R^2$

wherein X^1 is phenyl alanine

X^2 is any amino acid residue,

R¹ is NH₂- or an amino acid sequence X³-X⁴-X⁵

wherein X³ is an aliphatic amino acid residue having a side chain hydroxyl group
and X⁴ and X⁵ are the same or different and are any amino acid residue and wherein R² is a
sequence of 1 to 3 amino acids residues which are the same or different and are aliphatic
amino acid residues or of an effective fragment or derivative of said peptide.

14. (Amended) A method for treating or preventing anaphylactic hypotension in
a mammal comprising administering to the mammal an effective amount of a peptide of
[claim 1] the formula: R¹-X¹-X²-R²

wherein X¹ is phenyl alanine

X² is any amino acid residue,

R¹ is NH₂- or an amino acid sequence X³-X⁴-X⁵

wherein X³ is an aliphatic amino acid residue having a side chain hydroxyl group
and X⁴ and X⁵ are the same or different and are any amino acid residue and wherein R² is a
sequence of 1 to 3 amino acids residues which are the same or different and are aliphatic
amino acid residues or of an effective fragment or derivative of said peptide.

15. (Amended) A method of reducing or preventing an anaphylactic reaction in
a mammal comprising administering an effective amount of a peptide of [claim 1] the
formula: R¹-X¹-X²-R²

wherein X¹ is phenyl alanine

D3
 X^2 is any amino acid residue,

R^1 is NH_2 - or an amino acid sequence $X^3-X^4-X^5$

wherein X^3 is an aliphatic amino acid residue having a side chain hydroxyl group
and X^4 and X^5 are the same or different and are any amino acid residue and wherein R^2 is a
sequence of 1 to 3 amino acids residues which are the same or different and are aliphatic
amino acid residues or of an effective fragment or derivative of said peptide.

Please add new claims 22-30 as follows:

D4
--22. The peptide of claim 6 wherein at least one amino acid is a D amino acid.

23. A pharmaceutical composition comprising the peptide of claim 2 and a pharmaceutically acceptable carrier.

24. A pharmaceutical composition comprising the peptide of claim 6 and a pharmaceutically acceptable carrier.

SHELL
25. A method for treating or preventing SIRS-induced hypotension in a mammal comprising administering to the mammal an effective amount of the peptide of claim 2 or an effective fragment or derivative of said peptide.

26. A method for treating or preventing SIRS-induced hypotension in a mammal comprising administering to the mammal an effective amount of the peptide of claim 6 or an effective fragment or derivative of said peptide.

27. A method for treating or preventing anaphylactic hypotension in a mammal comprising administering to the mammal an effective amount of the peptide of claim 2 or an effective fragment or derivative of said peptide.

28. A method for treating or preventing anaphylactic hypotension in a mammal comprising administering to the mammal an effective amount of the peptide of claim 6 or an effective fragment or derivative of said peptide.

29. A method for treating or preventing an anaphylactic reaction in a mammal comprising administering to the mammal an effective amount of the peptide of claim 2 or an effective fragment or derivative of said peptide.

30. A method for treating or preventing an anaphylactic reaction in a mammal comprising administering to the mammal an effective amount of the peptide of claim 6 or an effective fragment or derivative of said peptide.--